CLAIM AMENDMENTS

- 1. (Original) Use of a biological photoreceptor as a light-controlled ion channel for the alteration of the ion conductivity of a membrane with the aid of light, wherein the photoreceptor used comprises an apoprotein and a light-sensitive polyene covalently bound to the apoprotein, said polyene interacting with the apoprotein and functioning as a light-sensitive gate.
- 2. (Original) Use according to Claim 1, characterised in that the apoprotein is a transmembrane protein with 5 or more transmembrane helices.
- 3. (Currently Amended) Use according to Claim 1 or 2, characterised in that the ion transport system is a proton transport system.
- 4. (Currently Amended) Use according to one of Claims 1 to 3 Claim 1, characterised in that the apoprotein is an opsin protein or a derivative or fragment of a naturally occurring opsin protein.
- 5. (Original) Use according to Claim 4, characterised in that the opsin derivative or fragment is the result of an exchange and/or an insertion and/or deletion of one or several amino acid(s) in the natural amino acid sequence of the opsin protein.
- 6. (Currently Amended) Use according to one of Claims 1 to 5 Claim 1, characterised in that the amino acid corresponding to the bacteriorhodopsin Asp⁹⁶ is an amino acid other than Asp and in the apoprotein at least 8 of the other 16 amino acids which are involved in the proton transport network in bacteriorhodopsin are identically retained or modified by conservative exchange.
- 7. (Currently Amended) Use according to one of Claims 1 to 6 Claim 1, characterised in that at least the amino acids which in bacteriorhodopsin correspond to the amino acids T⁴⁶, Y⁵⁷, R⁸², T⁸⁹, T¹⁰⁷, W¹⁸², D²¹² and K²¹⁶ are identically retained at the corresponding position.

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- 8. (Currently Amended) Use according to one of Claims 1 to 7 Claim 1, characterised in that the apoprotein contains the consensus sequence L(I)DxxxKxxW(F,Y).
- 9. (Currently Amended) Use according to one of Claims 1 to 8 Claim 1, characterised in that the apoprotein derives from lower plants.
- 10. (Original) Use according to Claim 9, characterised in that the lower plants are algae.
- 11. (Original) Use according to Claim 10, characterised in that the apoprotein is an opsin protein from *Chlamydomonas reinhardtii*.
- 12. (Currently Amended) Use according to one of Claims 1 to 11 Claim 1, characterised in that the apoprotein includes at least the amino acids 61 to 310 of the Channelopsin1 (CHOP-1) according to SEQ ID NO:AF385748 (National Center for Biotechnology Information, NCBI).
- 13. (Currently Amended) Use according to one of Claims 1 to 11 Claim 1, characterised in that the apoprotein includes at least the amino acids 24 to 268 of the Channelopsin2 (CHOP-2) according to SEQ ID NO:AF461397.
- 14. (Original) Use according to Claim 13, characterised in that the amino acid histidine at position 134 of the Channelopsin2 according to SEQ ID NO:AF461397 is replaced by another amino acid.
- 15. (Original) Use according to Claim 14, characterised in that the amino acid histidine at position 134 is replaced by arginine.
- 16. (Currently Amended) Use according to one of Claims 4 to 8 Claim 4, characterised in that the opsin protein derives from protozoa.

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- 17. (Currently Amended) Use according to one of Claims 4 to 8 Claim 4, characterised in that the opsin protein derives from bacteria or archaea.
- 18. (Currently Amended) Use according to one of Claims 4 to 8 Claim 4, characterised in that the opsin protein derives from fungi.
- 19. (Currently Amended) Use according to one of Claims 1 to 18 Claim 1, characterised in that the light-sensitive polyene is a retinal or retinal derivative.
- 20. (Original) Use according to Claim 19, characterised in that the retinal derivative is selected from the following group: 3,4-dehydroretinal, 13-ethylretinal, 9-dm-retinal, 3-hydroxyretinal, 4-hydroxyretinal, naphthylretinal; 3,7,11-trimethyl-dodeca-2,4,6,8,10-pentaenal; 3,7-dimethyl-deca-2,4,6,8-tetraenal; 3,7-dimethyl-octa-2,4,6-trienal; and 6-7 or 8-9 or 10-11 rotation-blocked retinals.
- 21. (Currently Amended) Use according to one of Claims 1 to 20 Claim 1 for the light-controlled alteration of the proton conductivity of the membrane.
- 22. (Currently Amended) Use according to one of Claims 1 to 20 Claim 1 for the light-controlled alteration of the membrane potential of a cell.
- 23. (Currently Amended) Use according to one of Claims 20 to 22 Claim 20, characterised in that the membrane is the cell membrane of a yeast, e.g. Saccharomyces cerevisiae, Schizosaccharomyces pombe or Pichia pastoris.
- 24. (Currently Amended) Use according to one of Claims 20 to 22 Claim 20, characterised in that the membrane is the cell membrane of a mammalian cell or insect cell, e.g. COS, BHK, HEK293, CHO, myeloma cell, MDCK or baculovirus-infected sf9 cell.
- 25. (Currently Amended) Use according to one of Claims 20 to 24 Claim 20 for the light-controlled raising or lowering of the intracellular concentration of ions.

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- 26. (Original) Use according to Claim 25 for the light-controlled raising or lowering of the intracellular proton concentration.
- 27. (Currently Amended) Use according to one of Claims 1 to 20 Claim 1 for the measurement of the intracellular proton concentration directly on the plasma membrane or of a proton concentration gradient across the plasma membrane with the aid of current-voltage curves, wherein the proton concentration gradient can be directly determined from the difference in the current-voltage curves with and without illumination from the reversal potential.
- 28. (Currently Amended) Use of a light-controlled ion channel according to one of Claims 1 to 20 Claim 1 for the high throughput screening of biological molecules.
- 29. (Original) Use according to Claim 28 for the high throughput screening of pH-regulated membrane proteins.
- 30. (Original) Use according to Claim 28 for the high throughput screening of voltage-dependent membrane proteins.
- 31. (Currently Amended) Use according to one of Claims 20 to 30 Claim 20, characterised in that the light-controlled ion channel is used in combination with a light-controlled active ion transport system.